## **Claims**

10

- 1. A tablet, comprising:-
  - (i) a core containing sumatriptan, and
- 5 (ii) a mantle, free of sumatriptan, wherein the mantle entirely surrounds the core.
  - 2. A tablet according to Claim 1, wherein the weight ratio of mantle:core is equal to or less than 1.8:1.
- 3. A tablet according to Claim 1, wherein the weight ratio of mantle:core is equal to or less than 1.5:1.
- 4. A tablet according to any of Claims 1-3, wherein the core contains from
  15 10-200 mg of sumatriptan.
  - 5. A tablet according to any of Claims 1-4, wherein:-
    - (i) the core is composed of sumatriptan, a filler, a binder, a disintegrant and a lubricant, and
- 20 (ii) the mantle is composed of a filler, a binder, a disintegrant and a lubricant.
  - 6. A tablet according to Claim 5, wherein the core and the mantle further comprise adsorbants and/or colorants.
  - 7. A tablet according to Claim 6, wherein the core comprises, by weight:-

sumatriptan: 1-40%

filler: 10-90%

binder: 2-60%

disintegrant: 1-60%

lubricant: 0.1-10%

adsorbants: 0-5%

colorants: 0-5%

25

30

and the mantle comprises, by weight:-

filler:

10-90%

binder:

2-60%

5

disintegrant: 1-60%

lubricant:

0.1-10%

adsorbants: 0-5%

colorants:

0-5%

A tablet according to Claim 6, wherein the core comprises by weight:-10 8.

sumatriptan

1-50%

filler:

10-90%

binder:

2-60%

15

disintegrant: 1-60%

lubricant:

0.1-10%

adsorbants:

0-5%

colorants:

0-5%

20 and the mantle comprises, by weight:-

filler:

10-90%

binder:

2-60%

disintegrant: 1-60%

25

lubricant:

0.1-10%

adsorbants: 0-5%

colorants:

0-5%

9. A tablet according to Claim 6, wherein the core comprises by weight:-

30

sumatriptan 5-80%

filler:

10-90%

binder:

2-60%

disintegrant: 1-60%

35

lubricant:

0.1-10%

adsorbants: 0-5% colorants: 0-5%

and the mantle comprises, by weight:-

5

filler:

10-90%

binder:

2-60%

disintegrant: 1-60%

lubricant:

0.1-10%

10 adsorbants: 0-5%

colorants:

0-5%

- A tablet according to any previous claim, wherein, apart from the 10. sumatriptan in the core, the core and the mantle are composed of substantially the same materials.
- A tablet according to any previous claim, wherein both the core and the 11. mantle dissolve rapidly in the stomach.
- 20 12. A tablet according to Claim 11, wherein at least 90% of the tablet is dissolved after 10 minutes.
  - 13. A tablet according to any of Claims 1-12, wherein the core and the mantle disintegrate over substantially the same time period.

25

35

15

- 14. A tablet according to Claim 13, wherein the mantle is at least 95% dissolved and the core is at least 90% dissolved after 10 minutes.
- A method of producing a tablet according to any previous claim, 15. 30 comprising the steps of:-
  - (a) forming a core by:-
    - (i) placing a first amount of powder/granule in a press.
    - compressing said first amount of powder/granule to obtain a (ii) core, and

10

15

- (b) pressing a second amount of powder/granule around said core, thereby forming a mantle and obtaining the final tablet.
- 16. A method of producing a tablet according to Claim 15, comprising the5 steps of:-
  - (a) forming a core by:-
    - (i) placing a first amount of powder/granule in a press,
    - (ii) compressing said first amount of powder/granule to obtain a core, and
  - (b) forming a mantle around the core by:-
    - (i) placing a second amount of powder/granule in a press,
    - (ii) placing said core onto said second amount of powder/granule,
    - (iii) placing a third amount of powder/granule on top of the core and the second amount of powder/granule, and
    - (iv) compressing (iii) so as to obtain the final tablet.
- 17. A method according to Claim 15 or 16, wherein the compression in Step (a) is carried out at pressure of from 0.5-5 tons.
  - 18. A method according to Claim 15 or 16, wherein the compression in Step (b) is carried out at a pressure from 0.5-10 tons.
- 25 19. A method according to Claim 15 or 16, wherein the first amount of powder/granule comprises sumatriptan, a filler, a binder, a disintegrant and a lubricant.
- 20. A method according to Claim 19, wherein the first amount of powder/granule further comprises an adsorbant and/or a colorant.
  - 21. A method according to Claim 15 or 16, wherein the first amount of powder/granule comprises, by weight:-
- 35 sumatriptan: 1-40%

filler:

10-90%

binder:

2-60%

disintegrant: 1-60%

lubricant:

0.1-10%

5 adsorbants: 0-5%

colorants:

0-5%

22. A method according to Claim 15 or 16, wherein the first amount of powder/granule comprises, by weight:-

10

sumatriptan 1-50%

filler:

10-90%

binder:

2-60%

disintegrant: 1-60%

15

;

lubricant:

0.1-10%

adsorbants:

0-5%

colorants:

0-5%

23. A method according to Claim 15 or 16, wherein the first amount of 20 powder/granule comprises, by weight:-

sumatriptan

5-80%

filler:

10-90%

binder:

2-60%

25 disintegrant: 1-60%

lubricant:

0.1-10%

adsorbants:

0-5%

colorants:

0-5%

- 30 24. A method according to Claim 15 or 16, wherein the second and/or third amounts of powder/granule comprise a filler, a binder, a disintegrant and a lubricant.
- A method according to Claim 24, wherein the second and/or third amounts 25. 35 of powder/granule further comprise an adsorbant and/or a colorant.

A method according to Claim 15 or 16, wherein the second and/or third 26. amounts of powder/granule comprise, by weight:-

filler:

10-90%

5

binder:

2-60%

disintegrant: 1-60%

lubricant:

0.1-10%

adsorbants:

0-5%

colorants:

0-5%

10

15

A method according to Claim 15 or 16, wherein Step (a) results in a 27. partially-compressed core, which core is then further compressed in Step (b).